Form PTO-1449 (REV. 7-80) U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

LIST OF PRIOR ART CITED BY APPLICANT

(Use several sheets if necessary)

Atty. Docket No. (Optional)	Application Number
17719 (PC27263A)	10/522,254

Applicant(s)
Manuela Villa, et al.

Filing Date	Group Art Unit
January 25, 2005	Unassigned

U.S. PATENT DOCUMENTS

EXAMINER INITIAL*		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (If appropriate)
	AA						
	AB						

U.S. PATENT APPLICATION PUBLICATIONS

EXAMINER INITIAL*		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (If appropriate)
VL	AA	2003/0073672 A1	4/17/03	Breitenbucher et al.			
	AB						
	AC						

FOREIGN PATENT DOCUMENTS

	REF	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES	TRANSLATION NO
VL		WO 02/064574 A2	8/22/02	PCT				
VL		WO 02/12242 A2	2/14/02	PCT				
VL		WO 96/12720	5/2/96	PCT				

OTHER DOCUMENTS *(Including Author, Title, Date, Pertinent Pages, Etc.)*

VL		Kikuchi C. et al., "Tetrahydrothienopyridylbutyl-Tetrahydrobenzindoles: New Selective Ligands of the 5-HT_7 Receptor", *Bioorganic & Medicinal Chemistry Letters, 12(18)*:2549-2552 (2002), XP-002256018
VL		Singh P. et al., "Quantitative Structure-Activity Relationship Studies on a New Class of Antihypertensive Agents: Derivatives of 3-Aryl-4,5,6,7-Tetrahydro-1H-Pyrazolo[4,3-c]Pyridine", *Quantitative Structure-Activity Relationships, 9(1)*:29-32 (1990), XP-001155089
VL		Winters G. et al., "Synthesis, in Vitro [^{3}H]Prazosin Displacement, and in Vivo Activity of 3-Aryl-4,5,6,7-Tetrahydropyrazolo[4,3-c]Pyridines, a New Class of Antihypertensive Agents", *J. Med. Chem., 28(7)*:934-940 (1985), XP-002256019

EXAMINER	DATE CONSIDERED
VL	12-07-06

* EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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1		Radinov R. et al., "3-Phenylpyrazolo[4,3-c]Pyridine and Derivatives: Structure Determination", *Journal of Molecular Structure, 158*:99-108 (1987), XP-009018257
2		Lackey K. et al., "The Discovery of Potent cRaf1 Kinase Inhibitors", *Bioorganic & Medicinal Chemistry Letters, 10(3)*:223-226 (2000), XP-004188821
3		Philip Cohen, "The Development and Therapeutic Potential of Protein Kinase Inhibitors", *Current Opinion in Chemical* Biology, *3(4)*:459-465 (1999), XP-002216616

EXAMINER	DATE CONSIDERED
	12-04-06

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